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NEWS 6 Apr 22 Records from IP.com available in CAPLUS, HCAPLUS, and
ZCAPLUS
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NEWS 8 Apr 22 Federal Research in Progress (FEDRIP) now available
NEWS 9 Jun 03 New e-mail delivery for search results now available
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NEWS 11 Jun 10 PCTFULL has been reloaded
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saved answer sets no longer valid
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NEWS 18 Aug 08 NTIS has been reloaded and enhanced
NEWS 19 Aug 19 Aquatic Toxicity Information Retrieval (AQUIRE)
now available on STN
NEWS 20 Aug 19 IFIPAT, IFICDB, and IFIUDB have been reloaded
NEWS 21 Aug 19 The MEDLINE file segment of TOXCENTER has been reloaded
NEWS 22 Aug 26 Sequence searching in REGISTRY enhanced
NEWS 23 Sep 03 JAPIO has been reloaded and enhanced
NEWS 24 Sep 16 Experimental properties added to the REGISTRY file
NEWS 25 Sep 16 CA Section Thesaurus available in CAPLUS and CA
NEWS 26 Oct 01 CASREACT Enriched with Reactions from 1907 to 1985
NEWS 27 Oct 21 EVENTLINE has been reloaded
NEWS 28 Oct 24 BEILSTEIN adds new search fields
NEWS 29 Oct 24 Nutraceuticals International (NUTRACEUT) now available on
STN
NEWS 30 Oct 25 MEDLINE SDI run of October 8, 2002
NEWS 31 Nov 18 DKLIT has been renamed APOLLIT
NEWS 32 Nov 25 More calculated properties added to REGISTRY
NEWS 33 Dec 02 TIBKAT will be removed from STN
NEWS 34 Dec 04 CSA files on STN
NEWS 35 Dec 17 PCTFULL now covers WP/PCT Applications from 1978 to date
NEWS 36 Dec 17 TOXCENTER enhanced with additional content
NEWS 37 Dec 17 Adis Clinical Trials Insight now available on STN
NEWS 38 Dec 30 ISMEC no longer available

NEWS 39 Jan 13 Indexing added to some pre-1967 records in CA/CAPLUS
NEWS 40 Jan 21 NUTRACEUT offering one free connect hour in February 2003
NEWS 41 Jan 21 PHARMAML offering one free connect hour in February 2003
NEWS 42 Jan 29 Simultaneous left and right truncation added to COMPENDEX,
ENERGY, INSPEC
NEWS 43 Feb 13 CANCERLIT is no longer being updated
NEWS 44 Feb 24 METADEX enhancements
NEWS 45 Feb 24 PCTGEN now available on STN
NEWS 46 Feb 24 TEMA now available on STN
NEWS 47 Feb 26 NTIS now allows simultaneous left and right truncation
NEWS 48 Feb 26 PCTFULL now contains images
NEWS 49 Mar 04 SDI PACKAGE for monthly delivery of multifile SDI results

NEWS EXPRESS January 6 CURRENT WINDOWS VERSION IS V6.01a,
CURRENT MACINTOSH VERSION IS V6.0b(ENG) AND V6.0Jb(JP),
AND CURRENT DISCOVER FILE IS DATED 01 OCTOBER 2002
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TOTAL
SESSION
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0.21
FULL ESTIMATED COST

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FILE COVERS 1907 - 13 Mar 2003 VOL 138 ISS 11
FILE LAST UPDATED: 12 Mar 2003 (20030312/ED)

This file contains CAS Registry Numbers for easy and accurate substance identification.

=> s 1,3-cyclic glycerophosphate
7465882 1
5712254 3
258238 CYCLIC
8095 GLYCEROPHOSPHATE
L1 2 1,3-CYCLIC GLYCEROPHOSPHATE
(1(W)3(W)CYCLIC(W)GLYCEROPHOSPHATE)

=> d 11 1-2 all

L1 ANSWER 1 OF 2 CAPLUS COPYRIGHT 2003 ACS
AN 2000:706969 CAPLUS
DN 133:261536
TI Pharmaceutical compositions comprising cyclic glycerophosphates and analogs thereof for promoting neural cell differentiation
IN Shinitzky, Meir
PA Yeda Research and Development Co. Ltd., Israel
SO PCT Int. Appl., 42 pp.
CODEN: PIXXD2
DT Patent
LA English
IC ICM A61K031-00
CC 1-11 (Pharmacology)
Section cross-reference(s): 29, 63

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2000057865	A2	20001005	WO 2000-IL185	20000324
	WO 2000057865	A3	20010628		
	W:	AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CR, CU, CZ, DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM			
	RW:	GH, GM, KE, LS, MW, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG			
	BR 2000009296	A	20011218	BR 2000-9296	20000324
	EP 1162959	A2	20011219	EP 2000-912877	20000324
	R:	AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO			
	JP 2002540146	T2	20021126	JP 2000-607616	20000324
PRAI	IL 1999-129178	A	19990325		
	WO 2000-IL185	W	20000324		
OS	MARPAT	133:261536			

AB Cyclic glycerophosphates and analogs thereof (CGs) are shown to exert neural promoting activities in target cells. Such activities include promotion of neuronal outgrowth, promotion of nerve growth, provision of dopaminotrophic supporting environment in a diseased portion of the brain,

prevention of nerve degeneration and nerve rescue. These activities of the CGs render them useful for treatment of various disorders including but not limited to mental disorders such as, for example, schizophrenia, dementia or disorders resulting in learning disabilities. In addn., these

CGs may be used for the treatment of neurodegenerative conditions such as Alzheimer's disease, Parkinson's disease, conditions resulting from exposure to harmful environmental factors or resulting from a mech. injury. The CGs may also be used to treat an individual suffering from a primary neurodegenerative condition in order to prevent or reduce the appearance of secondary degeneration in addnl. nerves ("nerve rescue"). For example, neural outgrowth of PC12 cells was seen when cells were

grown

in the presence of nerve growth factor (50 ng/mL) or 1,3
-cyclic glycerophosphate (1 .mu.M), but not in the presence of linear .alpha.-glycerophosphate.

ST cyclic glycerophosphate neuronal differentiation mental disorder; antipsychotic schizophrenia cyclic glycerophosphate; Alzheimer disease parkinsonism cyclic glycerophosphate

IT Anti-Alzheimer's agents
Antiparkinsonian agents
Antipsychotics
Mental disorder
Nervous system agents
Schizophrenia
(compns. comprising cyclic glycerophosphates for promoting neural differentiation for therapeutic uses)

IT Monoamines
Neurotrophic factors
RL: BPR (Biological process); BSU (Biological study, unclassified); BIOL (Biological study); PROC (Process)
(compns. comprising cyclic glycerophosphates for promoting neural differentiation for therapeutic uses)

IT Nerve
(degeneration, prevention of; compns. comprising cyclic glycerophosphates for promoting neural differentiation for therapeutic uses)

IT Mental disorder
(dementia; compns. comprising cyclic glycerophosphates for promoting neural differentiation for therapeutic uses)

IT Nerve
(differentiation; compns. comprising cyclic glycerophosphates for promoting neural differentiation for therapeutic uses)

IT Learning
(disorder; compns. comprising cyclic glycerophosphates for promoting neural differentiation for therapeutic uses)

IT Nerve
(dopaminergic, degeneration of; compns. comprising cyclic glycerophosphates for promoting neural differentiation for therapeutic uses)

IT Cell differentiation

(inducers; compns. comprising cyclic glycerophosphates for promoting neural differentiation for therapeutic uses)

IT Nerve, disease
(injury, neuronal rescue after; compns. comprising cyclic glycerophosphates for promoting neural differentiation for therapeutic uses)

IT Cell differentiation
Cell differentiation
(neuronal; compns. comprising cyclic glycerophosphates for promoting neural differentiation for therapeutic uses)

IT Drug delivery systems
(oral; compns. comprising cyclic glycerophosphates for promoting neural differentiation for therapeutic uses)

IT Drug delivery systems
(osmotic pumps; compns. comprising cyclic glycerophosphates for promoting neural differentiation for therapeutic uses)

IT Cell proliferation
(promotion of; compns. comprising cyclic glycerophosphates for promoting neural differentiation for therapeutic uses)

IT Drug delivery systems
(topical; compns. comprising cyclic glycerophosphates for promoting neural differentiation for therapeutic uses)

IT 298701-05-0P
RL: BAC (Biological activity or effector, except adverse); BPN (Biosynthetic preparation); BSU (Biological study, unclassified); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
(compns. comprising cyclic glycerophosphates for promoting neural differentiation for therapeutic uses)

IT 711-07-9P 13507-10-3P 22227-09-4P 118897-32-8P 123406-35-9P
286020-33-5P 298701-06-1P 298701-08-3P 298701-09-4P 298701-78-7P
RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
(compns. comprising cyclic glycerophosphates for promoting neural differentiation for therapeutic uses)

IT 51-61-6, Dopamine, biological studies 59-92-7, biological studies
102-32-9, DOPAC 306-08-1, Homovanillic acid
RL: BPR (Biological process); BSU (Biological study, unclassified); BIOL (Biological study); PROC (Process)
(compns. comprising cyclic glycerophosphates for promoting neural differentiation for therapeutic uses)

IT 9001-86-9, Phospholipase C
RL: BUU (Biological use, unclassified); BIOL (Biological study); USES (Uses)
(compns. comprising cyclic glycerophosphates for promoting neural differentiation for therapeutic uses)

IT 57-55-6, 1,2-Propanediol, reactions 96-26-4, Dihydroxyacetone
504-63-2, 1,3-Propanediol 770-12-7, Phenyl phosphorodichloridate
819-83-0, Disodium .beta.-glycerophosphate 4799-67-1 14690-00-7,
2-Benzylxy-1,3-propanediol 22002-87-5 26776-70-5, Dihydroxyacetone dimer
RL: RCT (Reactant); RACT (Reactant or reagent)
(compns. comprising cyclic glycerophosphates for promoting neural

differentiation for therapeutic uses)

IT 187976-16-5P
RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)
(compns. comprising cyclic glycerophosphates for promoting neural differentiation for therapeutic uses)

LL ANSWER 2 OF 2 CAPLUS COPYRIGHT 2003 ACS
AN 1993:534139 CAPLUS
DN 119:134139

TI Formation of 1,3-cyclic glycerophosphate by the action of phospholipase C on phosphatidylglycerol

AU Shinitzky, Meir; Friedman, Peter; Haimovitz, Rachel
CS Dep. Membrane Res. Biophys., Weizmann Inst. Sci, Rehovot, 76100, Israel
SO Journal of Biological Chemistry (1993), 268(19), 14109-15
CODEN: JBCHA3; ISSN: 0021-9258

DT Journal
LA English
CC 7-3 (Enzymes)

AB The action of phospholipase C (PLC) from *Bacillus cereus* on phosphatidylglycerol (PG), derived from egg yolk phosphatidylcholine (PC), was examd. in an ether-water mixt. The PLC cleavage of PG and PC followed a Michaelis-Menten kinetics with apparent V_{max} values per 1 μ g enzyme of 0.26 and 0.91 μ mol. min^{-1} and K_m values of 10 and 12 mM, resp. When the same enzymic reaction was carried out in minimally buffered aq. soln. of 1% Triton X-100, the decrease in pH with respect to phospholipid cleavage was as expected with PC but much less pronounced with PG. This could be accounted for by α -glycerophosphate, in the PLC hydrolysis of PG. Examn. of the chem. nature of the water-sol. product of PG by ^{31}P NMR revealed a single band at 2.31 ppm, while the bands of α -glycerophosphate and β -glycerophosphate appeared at 5.12 and 4.57 ppm, resp. Basic hydrolysis of the phospholipase cleavage product of PG (0.1 M NaOH for 1 min at 80 $^{\circ}$ C) followed by neutralization shifted its ^{31}P NMR band to 5.18 ppm, which practically coincided with that of α -glycerophosphate. Analogous expts. were carried out with PG labeled with 3H at the carbon 2 of the glycerol headgroup ($[^3H]PG$). Autoradiog. of thin layer chromatog. (TLC) of the $[^3H]PG$ enzymic hydrolyzate displayed a single 3H -labeled compd., which could be converted to α -glycerophosphate by basic hydrolysis. These results strongly suggest that the phosphate headgroup of PG is cleaved off by PLC as 1,3-cyclic glycerophosphate. A series of PLC expts. with phosphatidylhydroxyacetone and phosphatidyl 1,3-propanediol as model substrates supported this assignment. Two-dimensional homonuclear 1H NMR correlated spectra as well as IR spectra carried out on the isolated sodium salt of this product could further confirm such a structure. The unique structure and chem. nature of 1,3-cyclic glycerophosphate may bear a distinct physiol. function.

ST cyclic glycerophosphate formation phospholipase C phosphatidylglycerol
IT Phosphatidylglycerols
RL: RCT (Reactant); RACT (Reactant or reagent)

in) (cleavage of, by phospholipase C, cyclic glycerophosphate formation
IT Michaelis constant
 (of phospholipase C, with phosphatidylglycerol)
IT Phosphatidylcholines, reactions
 RL: RCT (Reactant); RACT (Reactant or reagent)
 (reaction of, with phospholipase C, kinetics of, phosphatidylglycerol
 in relation to)
IT 9001-86-9, Phospholipase C
 RL: BIOL (Biological study)
 (cyclic glycerophosphate formation by, of *Bacillus cereus*, in
 phosphatidylglycerol cleavage)
IT 42320-97-8
 RL: FORM (Formation, nonpreparative)
 (formation of, by phospholipase C cleavage of phosphatidylglycerol)
IT 149864-37-9
 RL: FORM (Formation, nonpreparative)
 (formation of, by phospholipase C cleavage of
 phosphatidylhydroxyacetone)
IT 13507-10-3
 RL: FORM (Formation, nonpreparative)
 (formation of, by phospholipase C cleavage of phosphatidylpropanediol)

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